## Amendments to the Claims

The listing of claims will replace all prior versions, and listings of claims in the application.

- 1. (currently amended) A method for obtaining human erythropoietin comprising
- (a) culturing mammalian cells which express recombinant human erythropoietin in culture medium consisting of:
  - (i) DMEM (Dulbecco's modified Eagle's medium) [5];
  - (ii) F12 medium [5];
  - (iii) insulin; and an additive, wherein the additive is selected from the group consisting of
  - (iv) NaHCO<sub>3</sub>, sugars, ethanolamine, pyruvate, <u>and</u> amino acids <u>as</u> additives and mixtures thereof,; and
  - (b) obtaining human erythropoietin from said culture medium.
- 2. (previously presented) The method of claim 1, wherein said cells are selected from the group consisting of CHO, COS, BHK, Namalwa, and HeLa.
  - 3. (original) The method of claim 2, wherein said cells comprise CHO cells.
- 4. (previously presented) The method of claim 1, wherein said culture medium comprises greater than about 1 mg insulin per liter of culture medium.

- 5. (previously presented) The method of claim 1, wherein said culture medium comprises less than about 20 mg insulin per liter of culture medium.
  - 6. (canceled)
- 7. (previously presented) The method of claim 1, wherein said obtaining comprises:
- (i) separating supernatant comprising EPO and insulin from said cells;
  - (ii) concentrating supernatant of step (i); and
  - (iii) freezing concentrated product of step (ii).
- 8. (previously presented) The method of claim 7, wherein media is added to separated cells of step (i) and said cells are cultured.
- 9. (previously presented) The method of claim 7, wherein supernatant of said step (i) is concentrated from about 50 to 150 fold.
- 10. (previously presented) The method of claim 7, wherein supernatant of said step (i) is concentrated about 100 fold.
- 11. (previously presented) The method of claim 7, wherein said step (ii) comprises using a tangential filtration system through membranes with a molecular weight cut-off of about 3,000 Daltons.

- 12. (previously presented) The method of claim 7, further comprising (iv) sterile filtering the concentrated product of step (iii) through membranes with pores of diameters of about  $0.2~\mu m$ .
- 13. (previously presented) The method of claim 1, wherein said culture medium comprises about 10 mg insulin per liter of culture medium.
  - 14. (canceled)
- 15. (previously presented) The method of claim 1, wherein said sugars are selected from the group consisting of glucose, lactose, galactose and mixtures thereof.
- 16. (previously presented) The method of claim 1, wherein said pyruvate is sodium pyruvate.
- 17. (previously presented) The method of claim 1, wherein said amino acids are selected from the group consisting of glutamine, tryptophan, asparagine, serine and mixtures thereof.
  - 18. (canceled)
- 19. (currently amended) The method of claim 1 18, wherein said culture medium contains Iscove's DMEM, HAM's F12 medium, insulin and NaHCO<sub>3</sub>, glucose,

lactose, galactose, ethanolamine, sodium pyruvate, glutamine, tryptophan, asparagine and serine as additives.

- 20. (previously presented) The method of claim 1, wherein said DMEM is Iscove's DMEM and wherein said F12 medium is HAM's F12 medium.
  - 21. (New) A method for obtaining human erythropoietin comprising
- (a) culturing mammalian cells which express recombinant human erythropoietin in culture medium consisting of:
  - (i) DMEM (Dulbecco's modified Eagle's medium);
  - (ii) F12 medium;
  - (iii) insulin; and
  - (iv) NaHCO<sub>3</sub>, glucose, lactose, galactose, ethanolamine, sodium pyruvate, glutamine, tryptophan, asparagine and serine as additives; and
  - (b) obtaining human erythropoietin from said culture medium.
  - 22. (New) The method of claim 21, wherein said cells are CHO cells.